UTAH DEPARTMENT OF HEALTH, PRIOR AUTHORIZATION REQUEST FORM

Verquvo (vericiguat)

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Member and Medication Information (required)			
Member ID:		Member Name:	
DOB:		Weight:	
Medication Name/ Strength:		Dose:	
Directions for use:			
Provider Information (required)			
Name:	NPI:		Specialty:
Contact Person:	Office Phone:		Office Fax:
FAX FORM AND RELEVANT DOCUMENTATION INCLUDING: LABORATORY RESULTS, CHART NOTES and/or UPDATED PROVIDER LETTER TO 855-828-4992			
Criteria for Approval (ALL of the following criteria must be met): 18 years of age or older. Not pregnant or breast feeding. Not taking soluble guanylate cyclase stimulators (e.g. riociguat). Not taking phosphodiesterase type 5 inhibitors (e.g. sildenafil, tadalafil, vardenafil). Diagnosed with chronic heart failure with an ejection fraction 45% or less who are NYHA class II-IV and either: Hospitalized due to heart failure within last 6 months. OR Required IV diuretics as an outpatient within the previous 3 months. Concurrently receiving the maximum tolerated or target dose of guideline-directed medical therapy for heart failure, unless not tolerated or contraindicated: Beta-blockers (carvedilol, metoprolol succinate, or bisoprolol). Medication and dose: Angiotensin antagonist (ARNI, ACEI, ARB) Medication and dose: Mineralocorticoid receptor antagonist (e.g. spironolactone) if LVEF < 35% or LVEF ≤ 40% with diabetes mellitus or post myocardial infarction with HF symptoms. Medication and dose: ———————————————————————————————————			
Note: Medication has not been studied in patients with end-stage-renal disease or hepatic insufficiency. Recommend monitoring according to package insert.			
Re-authorization Criteria: Updated letter with medical justification or updated chart notes demonstrating positive clinical response.			
Initial Authorization: Up to six (6) months Re-authorization: Up to one (1) year			
PROVIDER CERTIFICATION			
I hereby certify this treatment is indicated, necessary and meets the guidelines for use.			
Prescriber's Signature		 Date	